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PATENT

ATTORNEY DOCKET NO.: 056291-5231

IN THE UNITED STATES PATENT AND TRADEMARK OFFICE

In re Application of:

Andrew CASSIDY et al.

Confirmation No.: 2301

Application No.: 10/568,432

Group Art Unit: 1645

Filed: February 14, 2006

Examiner: Unassigned

For: AMPLIFICATION METHOD

Commissioner for Patents
U.S. Patent and Trademark Office
Customer Window,
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Alexandria, VA 22314

Sir:

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There is an error with respect to the following data which is:

incorrectly entered

Error In

Correct Data

Domestic Priority Data

PCT/GB04/03486 August 12, 2004

Attached is a copy of the WO 2005/019452 A1 Cover Page

The correction is not due to any error by the applicants and no fee is due.

Application No.: 10/568,432 Attorney Docket No.: 056291-5231

The above correction is due to the applicants error and the fee therefor under 37 CFR 1.19(h) of \$25.00 is paid as follows:

The Commissioner is hereby authorized to charge any fees which may be required, or credit any overpayment to Deposit Account 50-0310.

Respectfully submitted,

MORGAN, LEWIS & BOCKIUS LLP

for Sally B. Told

Reg. No: 45,397

Dated: February 20, 2007

MORGAN, LEWIS & BOCKIUS LLP

1111 Pennsylvania Ave, N.W. Washington, D.C. 20004 (202) 739-3000



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MORGAN LEWIS & BOCKIUS LLP 1111 PENNSYLVANIA AVENUE NW WASHINGTON, DC 20004

CONFIRMATION NO. 2301

FILING RECEIPT

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Date Mailed: 02/07/2007

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Applicant(s)

Andrew Cassidy, Cheshire, UNITED KINGDOM; Alexander Graham, Cheshire, UNITED KINGDOM;

Power of Attorney: The patent practitioners associated with Customer Number <u>09629</u>.

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The country code and number of your priority application, to be used for filing abroad under the Paris Convention, is **US10/568,432**

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MORGAN, LEWIS & BOCKIUS LLP

Title

Amplification method

Preliminary Class

435

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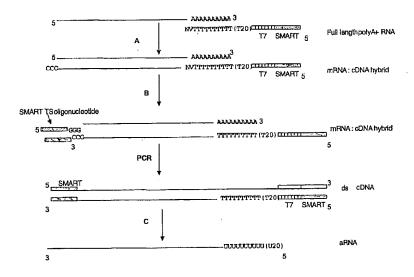
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(71) Applicant (for AE, AG, AL, AM, AT, AU, AZ, BA, BB, BE, BG, BR, BW, BY, BZ, CA, CH, CN, CO, CR, CU, CY, CZ, DE, DK, DM, DZ, EC, EE, EG, ES, FI, GB, GD, GE, GH, GM, HR, HU, ID, IL, IN, IS, JP, KE, KG, KP, KR, KZ, LC, LK, LR, LS, LT, LU, LV, MA, MD, MK, MN, MW, MX, MZ, NA, NI, NO, NZ, OM, PG, PH, PL, PT, RO, RU, SC, SD, SE, SG, SK, SL, SY, SZ, TJ, TM, TN, TR, TT, TZ, UA, UG, UZ, VC, VN, YU, ZA, ZM, ZW only): ASTRAZENECA AB [SE/SE]; Sodertalje, S-151 85 (SE).

- (71) Applicant (for MG only): ASTRAZENECA UK LIM-ITED [GB/GB]; 15 Stanhope Gate, London Greater London W1K 1LN (GB).
- (72) Inventors; and
- (75) Inventors/Applicants (for US only): GRAHAM, Alexander [GB/GB]; AstraZeneca R & D Alderley, Alderley Park, Macclesfield Cheshire SK10 4TG (GB). CASSIDY, Andrew [GB/GB]; AstraZeneca R & D Alderley, Alderley Park, Macclesfield Cheshire SK10 4TG (GB).
- (74) Agent: ASTRAZENECA; Global Intellectual Property, S-151 85 Sodertalje (SE).
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[Continued on next page]

(54) Title: AMPLIFICATION METHOD



(57) Abstract: The present invention combines the use of template switching with nucleic acid amplification - such as PCR - to generate amplification products representative of an entire RNA population. An RNA polymerase promoter sequence allows transcription-based amplification to be performed on the derived amplification products such that antisense amplified RNA (aRNA) or complementary RNA (cRNA) is produced for subsequent downstream applications.

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